ACID REDUCER MAXIMUM STRENGTH- famotidine tablet, film coated Rite Aid Corporation

Rite Aid Corporation Acid Reducer Drug Facts

Active ingredient (in each tablet)

Famotidine 20 mg

Purpose

Acid reducer

Uses

- relieves heartburn associated with acid indigestion and sour stomach
- prevents heartburn associated with acid indigestion and sour stomach brought on by eating or drinking certain food and beverages

Warnings

Allergy alert: Do not use if you are allergic to famotidine or other acid reducers

Do not use

- if you have trouble or pain swallowing food, vomiting with blood, or bloody or black stools. These may be signs of a serious condition. See your doctor.
- with other acid reducers

Ask a doctor before use if you have

- had heartburn over 3 months. This may be a sign of a more serious condition.
- heartburn with **lightheadedness**, **sweating**, **or dizziness**
- chest pain or shoulder pain with shortness of breath; sweating; pain spreading to arms, neck or shoulders; or lightheadedness
- frequent **chest pain**
- frequent wheezing, particularly with heartburn
- unexplained weight loss
- nausea or vomiting
- stomach pain
- kidney disease

Ask a doctor or pharmacist before use if you are

taking a prescription drug. Acid reducers may interact with certain prescription drugs.

Stop use and ask a doctor if

- your heartburn continues or worsens
- you need to take this product for more than 14 days

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

- adults and children 12 years and over:
- to **relieve** symptoms, swallow 1 tablet with a glass of water. Do not chew.
- to **prevent** symptoms, swallow 1 tablet with a glass of water at any time from **10 to 60 minutes before** eating food or drinking beverages that cause heartburn
- do not use more than 2 tablets in 24 hours
- children under 12 years: ask a doctor

Other information

- read the directions and warnings before use
- keep the carton. It contains important information.
- store at 20°-25°C (68°-77°F)
- protect from moisture and light

Inactive ingredients

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, talc, titanium dioxide

Questions or comments?

1-800-719-9260

Principal Display Panel

SEE NEW WARNINGS

Compare to the active ingredient of Maximum Strength Pepcid® AC

ACID REDUCER

Famotidine Tablets, 20 mg

acid reducer

MAXIMUM STRENGTH

Just one tablet prevents & relieves heartburn due to acid indigestion

ACTUAL SIZE

25 TABLETS



ACID REDUCER

Famotidine Tablets, 20 mg

acid reducer

MAXIMUM STRENGTH

Just one tablet prevents & relieves heartburn due to acid indigestion

ACTUAL SIZE



ACID REDUCER MAXIMUM STRENGTH

famotidine tablet, film coated

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Product TypeHUMAN OTC DRUGItem Code (Source)NDC:11822-0194

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FAMO TIDINE (UNII: 5QZO15J2Z8) (FAMOTIDINE - UNII:5QZO15J2Z8)	FAMOTIDINE	20 mg

Inactive Ingredients				
Ingredient Name	Strength			
CARNAUBA WAX (UNII: R12CBM0EIZ)				
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)				
CROSCARMELLOSE SODIUM (UNII: M28 O L1HH48)				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)				
TALC (UNII: 7SEV7J4R1U)				
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)				

Product Characteristics						
Color	WHITE	Score	no score			
Shape	ROUND	Size	8mm			
Flavor		Imprint Code	L194			
Contains						

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11822-0194-3	1 in 1 CARTON	09/29/2006		
1		50 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:11822-0194-5	25 in 1 CARTON	09/29/2006		
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product			
3	NDC:11822-0194-7	2 in 1 CARTON	09/29/2006		
3		85 in 1 BOTTLE; Type 0: Not a Combination Product			
4	NDC:11822-0194-2	30 in 1 CARTON	04/13/2008	07/29/2015	

4 1	1 in 1 BLISTER PACK; Type 0: Not a Combination Product					
Marketing Information						
Wai keung inioi mauon						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
ANDA	ANDA077351	09/29/2006				

Labeler - Rite Aid Corporation (014578892)

Revised: 7/2020 Rite Aid Corporation